Heritability and Tissue Specificity of Expression Quantitative Trait Loci

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Variation in gene expression is heritable and has been mapped to the genome in humans and model organisms as expression quantitative trait loci (eQTLs). We applied integrated genome-wide expression profiling and linkage analysis to the regulation of gene expression in fat, kidney, adrenal, and heart tissues using the BXH/HXB panel of rat recombinant inbred strains. Here, we report the influence of heritability and allelic effect of the quantitative trait locus on detection of cis- and trans-acting eQTLs and discuss how these factors operate in a tissue-specific context. We identified several hundred major eQTLs in each tissue and found that cis-acting eQTLs are highly heritable and easier to detect than trans-eQTLs. The proportion of heritable expression traits was similar in all tissues; however, heritability alone was not a reliable predictor of whether an eQTL will be detected. We empirically show how the use of heritability as a filter reduces the ability to discover trans-eQTLs, particularly for eQTLs with small effects. Only 3% of cis- and trans-eQTLs exhibited large allelic effects, explaining more than 40% of the phenotypic variance, suggestive of a highly polygenic control of gene expression. Power calculations indicated that, across tissues, minor differences in genetic effects are expected to have a significant impact on detection of trans-eQTLs. Trans-eQTLs generally show smaller effects than cis-eQTLs and have a higher false discovery rate, particularly in more heterogeneous tissues, suggesting that small biological variability, likely relating to tissue composition, may influence detection of trans-eQTLs in this system. We delineate the effects of genetic architecture on variation in gene expression and show the sensitivity of this experimental design to tissue sampling variability in large-scale eQTL studies.


Introduction

Quantitative variation in gene expression levels acts as an intermediate phenotype situated between genomic DNA sequence variation and more complex cellular, organ, or whole body phenotypes. Numerous studies indicate that individual variation in gene expression (i.e., transcript abundance) isheritable in segregating populations [1–5]. Gene expression can therefore be mapped to the genome using standard linkage methods, allowing identification of expression quantitative trait loci (eQTLs), which represent genomic regions for the genetic control of gene expression [6]. This approach has been termed genetical genomics [7,8], and recent technological and methodological advances have made its large-scale application feasible at the level of the genome. One of the most powerful features of this approach is the ability to discriminate between cis- and trans-acting influences on gene expression and potentially to dissect complex regulatory networks [9,10]. A cis-acting eQTL maps to the physical location of the gene itself, whereas a trans-acting eQTL maps to a genomic region that is distant from the physical location of the gene being transcribed. By combining the genomic position of the gene encoding each transcript and the position of its eQTL, it is possible to discriminate between cis- and trans-regulatory control elements of gene expression for thousands of genes across the genome.

A number of genetical genomics studies show that sequence variation in cis-acting genes plays a considerable role in determining detectable variability in gene expression, and, accordingly, cis effects are usually mapped with high statistical significance [3,11]. cis-acting genes are generally easier to detect by linkage, since they explain a large fraction of the variance of gene expression, and are of great interest as positional candidates for physiological quantitative trait loci (QTLs) [12]. Although trans-eQTLs may be associated with lesser statistical significance, they are often detected as clusters, reflecting coordinated regulation of many genes by a single “master regulator” [5]. Trans-acting eQTL genes usually explain <20% of the phenotypic variance and may be below a stringent threshold of detection for linkage, since
Synopsis

The combined application of genome-wide expression profiling from microarray experiments with genetic linkage analysis enables the mapping of expression quantitative trait loci (eQTLs), which are primary control points for gene expression across the genome. This approach has been called "genetical genomics" and recent technological and methodological advances have made its large-scale application feasible in humans and model organisms. Using this approach, the authors have carried out an extensive analysis of the genetic architecture underlying variation in gene expression using a panel of 30 rat recombinant inbred strains. The results are used to explore the relationship between heritability of gene expression, cis- and trans-acting genetic effects, tissue heterogeneity, and statistical cut-offs of significance, which are important factors for large-scale eQTL studies. By examining large eQTL data from four tissues, the authors provide a detailed picture of cis- and trans-eQTL features that may help understanding of the genetic regulation of transcription on a genomic scale. The results also show the sensitivity of this approach to discriminate between cis and trans regulation and the value of the rat system in studying large eQTL datasets from multiple tissues.

Results

Heritability of Gene Expression in RI Strains

Gene expression profiles and trait heritability ($h_{\text{trait}}^2$) across 30 RI strains were investigated in fat, kidney, adrenal, and left ventricle (LV) tissue for each transcript considered in this study (see Materials and Methods). The median (25%–75% quartiles) $h_{\text{trait}}^2$ is 0.14 (0.11–0.18) for LV, 0.17 (0.13–0.22) for fat, 0.14 (0.10–0.20) for kidney, and 0.17 (0.12–0.24) for adrenal. We observed that the distribution of heritable gene expression is comparable across tissues ($p > 0.05$, chi-square test; Figure S1), suggesting a similar extent of the genetic component segregating in the RI strains. The estimates of $h_{\text{trait}}^2$ were positively correlated with the average levels of expression (Spearman's rank correlations: 0.33 in LV, 0.47 in fat, 0.1 in kidney, and 0.4 in adrenal, $p < 0.05$ in all tissues). We calculated the number of genes exhibiting different $h_{\text{trait}}^2$ and observed no significant difference in the proportions of heritable transcripts across four tissues (Figure S2).

Global Analysis of Heritability and Allelic Effect of eQTLs in Four Tissues

Table 1 summarises the mapping results in four tissues for the major eQTLs (i.e., the eQTL with the highest statistical significance for each probe set) detected in cis or trans for each transcript. The median for $h_{\text{cis}}^2$ QTL effects, and heritability of the eQTLs ($h_{\text{QTL}}^2$) are also reported for each threshold of significance. For all major eQTLs detected at $p = 0.05$, the median $h_{\text{trait}}^2$ ranges from a minimum of 0.14 for the trans-eQTLs to a maximum of 0.37 for the cis-eQTLs. At higher levels of significance the $h_{\text{trait}}^2$ increases for both cis- and trans-acting eQTLs, but it is consistently lower for the trans-eQTLs (Table 1).

At genome-wide significance ($p = 0.05$) eQTL effects are relatively small, ranging from 0.06 (trans-eQTLs) to 0.18 (cis-eQTLs). Median allelic effects increase at lower $p$-values up to 0.37 for cis-acting eQTLs and 0.11 for trans-acting eQTLs. The distribution of allelic effects in the range 0–1.5 (i.e., absolute FC in the range 1–8) and relative $h_{\text{QTL}}^2$ for all major eQTLs

Norway (BN) strain to generate the BXH/HXB panel of 30 RI lines [18], one of the most widely used model systems of human hypertension and metabolic syndrome [18–21]. Along with other features of the RI strains, the ability to make measurements in multiple genetically identical animals from the same strain increases trait heritability, thus facilitating eQTL identification [22,23]. This makes the RI lines an ideal model system to investigate the heritable component of gene expression for cis- and trans-acting regulators in the context of natural variation in complex physiological processes. Given that in similar study designs [16,17,24] high trait heritability has been used as one of the criteria for statistical significance to map eQTLs, we investigate the effect of heritability filtering on detection of cis- and trans-acting eQTLs. Here, we show how the overall sensitivity of the genetical genomics approach is determined by factors such as patterns of heritability of gene expression and tissue-specific genetic effects. We discuss how these factors affect the relative power to detect cis- and trans-eQTLs and highlight the importance of quantifying heritable determinants of gene expression in the context of single tissues.
mapped with genome-wide significance is reported in Figure 1. The vast majority of the eQTLs explained less than 40% of the genetic variance, suggesting that undetected eQTLs account for the rest of the variance in such transcripts. Notably, the strongest eQTLs explained >30% of genetic variance for >26% of the transcripts, whereas only ~3% of both of cis- and trans-acting eQTLs (detected at high significance levels, \( p < 10^{-5} \)) exhibited considerably large QTL effects and \( h^2_{\text{QTL}} > 0.4 \).

Heritability of Gene Expression for Detection of Cis- and Trans-eQTLs

We investigated the \( h^2_{\text{trait}} \) as a predictor of the existence of all significant eQTLs in LV, fat, kidney, and adrenal tissue, using receiver operating characteristic (ROC) curves (Figure S5). For the eQTLs detected at \( p = 0.05 \), the maximum area under the ROC curve (AUC) is 0.73 (standard error [SE] 0.007), which is commonly associated with inaccurate prediction [25]. Heritability appears to be a better predictor (minimum AUC in all tissues = 0.88 [SE 0.011], average AUC across tissues = 0.91) only for the eQTLs detected at \( p < 10^{-5} \) (Table S1). The ability of the \( h^2_{\text{trait}} \) to predict the existence of an eQTL improves by increasing the cut-off of significance for linkage, and at \( p = 10^{-4} \) the AUC is greater than 0.95 in all tissues (Figure S3; Table S1). We then considered the ability of heritability to predict the existence of cis- or trans-eQTLs separately. In all tissues, ROC analysis showed that \( h^2_{\text{trait}} \) is a poor predictor of the trans-eQTLs detected at \( p = 10^{-2} \). AUC ranges from 0.47 to 0.53. At each threshold of significance, the null hypothesis that the AUC equals 0.5 cannot be rejected for the trans-eQTLs. As such, the \( h^2_{\text{trait}} \) is no better than chance in predicting a detectable trans-eQTL. For the cis-eQTLs, we observed higher accuracy in the prediction: AUC ranges from 0.69 (eQTLs detected at \( p = 10^{-5} \)) to 0.77 (eQTLs detected at \( p = 10^{-5} \)), and the AUC is significantly different from 0.5 at each level of significance.

When high trait heritability was used as the criterion to identify transcripts for which genetic linkage is expected to be more reliable, a significant proportion of eQTLs could not be detected. Given the actual number of eQTLs mapped with genome-wide significance in this study, we calculated the percentage of cis- and trans-eQTLs that will be discarded when various \( h^2_{\text{trait}} \) cut-offs are used (Figure 2). For example, if \( h^2_{\text{trait}} > 20\% \) is accepted as a prerequisite to map an eQTL and hence eQTLs called from transcripts showing \( h^2_{\text{trait}} < 20\% \) are not considered, 65% to 80% of the trans-eQTLs and 10% to 40% of the cis-eQTLs will be excluded from the eQTL dataset (Figure 2).

Power to Detect Cis and Trans Effects and False Discovery Rate

Figure 3A shows the power for the minimum detectable effect at different \( h^2_{\text{trait}} \) values when allelic effects are in the range 0–0.30 (i.e., absolute FC between one and 1.52). For effects greater than 0.25, the power tends to 100% for all considered values of \( h^2_{\text{trait}} \). Small effects are detected with low power, and when the \( h^2_{\text{trait}} \) is 0.8–0.9, we achieve ~75% power for an effect of 0.10, which is equivalent to an absolute FC of 1.15. For lower heritabilities of the trait (\( h^2_{\text{trait}} = 0.2–0.3 \)), the minimum detectable eQTL effect is higher, ~0.12 (i.e., an absolute FC of 1.18). The power to detect small effects increases with the number of biological replicates within each strain, with a significant improvement observed in the range of two to six replicates (Figure 4), and with the number of KI lines (Figure S5). Similar analyses showed that the power increases for eQTLs mapped at higher significance levels (unpublished data), indicating that stronger effects are associated with lower \( p \)-values.

Given the effects for cis- and trans-eQTLs detected with
genome-wide significance in this study (Table 1), we derived the expected power for the cis and trans allelic effects in the LV, fat, kidney, and adrenal tissue. Figure 3B shows that the maximum expected power to detect average effects of trans-eQTLs is low in all tissues, ranging from 12% to 60% (at high $h^2_{trait}$). In contrast, the cis-regulated genes are expected to be detectable with high power across all tissues. As indicated by tissue-specific allelic effects of the trans-eQTL, we expect different power for trans-regulated genes across tissues (Figure 3B).

The lower power to detect trans-eQTLs at $p = 0.05$ in kidney and adrenal is reflected in the tissue-specific false discovery rate (FDR) (Figure 4A). At $p = 0.05$, we observe an FDR of $\sim 35\%$ in kidney and adrenal, whereas the FDR is only 26% in LV and fat. This difference in FDR is noticeable for a $p$-value cut-off of $>10^{-5}$, and below this threshold all linkage datasets show similar FDR patterns (Figure 4A, insert). Given that trans-acting eQTLs are normally detected at higher $p$-value thresholds, they show higher FDR levels than cis-regulated genes (Figure 4B). Across all tissues, at $p = 0.05$ the median FDR ranges from 4% to 8% for the cis-eQTLs and from 19% to 27% for the trans-eQTLs. For trans-acting eQTLs, kidney and adrenal show higher FDR levels (median 24%–27%) than LV and fat tissues (median FDR 19%–20%).

**Discussion**

We investigated the overall sensitivity of the genetical genomics approach to discriminate between cis and trans regulation within and between tissues. Our analysis was carried out in the BXH/HXB panel of rat RI strains and in four tissues: LV, fat, kidney, and adrenal, from which expression profiles were generated. RI strains are a suitable genetic system for global analysis of heritable patterns of gene expression, allowing direct estimation of genetic and environmental components of phenotypic variance [22]. We examined how cis and trans genetic factors contributed to the global heritability of gene expression observed across tissues. Quantifying the extent of such contributions is of great importance to understand how genetic influences of gene expression are structured within the population and may occur unequally in the context of specific tissues [26].

We provided evidence for a significant heritable component of quantitative variation of gene expression in all tissues. On the whole, at least 20% of the transcripts showed $h^2_{trait} > 20\%$, and the overall proportion of heritable genes was similar across all tissues (Figure S2). The value of $h^2_{trait}$ is higher for those transcripts that were mapped to cis-acting eQTLs (31%–51%) than trans-eQTLs (12%–21%), thus
suggested that larger differences in transcript abundance are associated with cis-acting genetic variation in gene expression (Table 1). Although $h^2_{\text{trait}}$ is positively correlated with the average levels of expression, the proportions of heritable transcripts do not vary by the levels of expression observed in different tissues (unpublished data). We also observed that amount of variation in gene expression levels is consistent across tissues, indicating that the estimates of heritability are not significantly influenced by intra-specific variability between tissues (see Materials and Methods).

It is commonly accepted that the power and precision of genetic mapping will be significantly affected by the magnitude of trait heritability [27]. In practice, the larger the environmental variance (i.e., lower $h^2_{\text{trait}}$), the less likely an eQTL will be detected. However, detection of a significant eQTL does not appear to be entirely determined by the sole estimate of $h^2_{\text{trait}}$. We showed that in large-scale analysis of gene expression phenotypes, $h^2_{\text{trait}}$ by itself is not a consistent predictor of the eQTLs, and its efficiency depends considerably on the $p$-value thresholds at which eQTLs are mapped (Figure S3; Table S1). In addition, we have shown that at each level of significance of the eQTL, $h^2_{\text{trait}}$ is an unreliable parameter in predicting putative trans-eQTLs. When $h^2_{\text{trait}}$ is used as a filter to improve detection of linkages [16,17,24], a significant proportion of eQTLs will be discarded from the study. For example, we have shown that conservative filtering approaches ($h^2_{\text{trait}} > 20\%$) result in a significant decrease (65%–80%) in the trans-acting eQTLs detected with genome-wide significance (Figure 2). This could have important consequences for identification of clusters of trans-acting eQTLs, or trans-eQTL “hotspots”, which may represent common regulation by a single gene. It should also be noted that the relative proportion of trans-acting eQTLs tends to drop as the statistical significance for eQTL detection is increased (Table 1). This suggests that using conservative filtering approaches may result in many false negatives and in fewer trans-eQTL “hotspots” being identified, biasing detection towards cis-acting eQTLs.

The relationship between $h^2_{\text{trait}}$ and different detection of cis- and trans-eQTLs may be a result of their specific mono-, oligo-, or polygenic influences on gene expression. Because sequence variations within single cis-acting genes may have a substantial impact on transcription, the measured heritability of gene expression can be very high, indicating existence of major cis-regulated eQTLs that explain most of the phenotypic variance [13]. This argument may account for the better efficiency of $h^2_{\text{trait}}$ in predicting highly significant cis-eQTLs, since most of them are expected to be inherited essentially as monogenic traits. We found that whilst cis-eQTLs have substantial $h^2_{\text{QTL}}$ (26%–31%), the median $h^2_{\text{QTL}}$ observed for trans-eQTLs mapped at $p = 0.05$ is only 20%–23%. However, at higher levels of significance we observed more robust $h^2_{\text{QTL}}$ values for both cis and trans loci (Table 1). This can be indicative of an over-estimate of the eQTL effect size due to the relatively small sample size (Beavis effect [28]) or of a combined effect of a number of physically linked QTLs of small effect [27]. Importantly, we detected only a small proportion of eQTLs (mostly cis-acting) that exhibit big allelic effects and account for more than 40% of the phenotypic variance. These data suggest that global variation in gene expression is also likely to be under the influence of multiple loci with small effect alongside a major cis-eQTL. This explanation is consistent with previous data in yeast, where only 3% of highly

Figure 2. Proportion of eQTLs That Are Excluded when Transcripts are Filtered Based on $h^2_{\text{trait}}$

Percentages were calculated as follows: $(1 - \text{eQTL}_{\text{filtered}}/\text{eQTL}_{\text{total}}) \times 100$, where eQTL$_{\text{total}}$ is the total number of eQTLs detected at $p = 0.05$ without filtering based on $h^2_{\text{trait}}$ (see Table 1), and eQTL$_{\text{filtered}}$ is the total number of eQTLs observed when transcripts were filtered based on a given trait heritability cut-off. For any trait heritability cut-off, the number of excluded eQTLs is higher for analysis of gene expression phenotypes, eQTLs. When $h^2_{\text{trait}}$ have shown that at each level of significance of the eQTL, the $h^2_{\text{trait}}$ of linkages [16,17,24], a significant proportion of eQTLs will be discarded from the study. For example, we have shown that conservative filtering approaches ($h^2_{\text{trait}} > 20\%$) result in a significant decrease (65%–80%) in the trans-acting eQTLs detected with genome-wide significance (Figure 2). This could have important consequences for identification of clusters of trans-acting eQTLs, or trans-eQTL “hotspots”, which may represent common regulation by a single gene. It should also be noted that the relative proportion of trans-acting eQTLs tends to drop as the statistical significance for eQTL detection is increased (Table 1). This suggests that using conservative filtering approaches may result in many false negatives and in fewer trans-eQTL “hotspots” being identified, biasing detection towards cis-acting eQTLs.

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heritable transcripts are explained by single-locus (i.e., monogenic) inheritance and 50% are consistent with more than five controlling loci of equal effect [29,30].

We and others [5] detected a substantial relative proportion (54%–64%) of trans-acting eQTLs at \( p = 0.05 \), and this percentage decreases at higher levels of significance [11]. At each cut-off for linkage, the median effect observed for trans-eQTLs (0.06–0.11, FC 1.09–1.16) is well below that observed for the cis-eQTLs (0.14–0.37, FC 1.21–1.67). This suggests that detection of cis-eQTLs will be facilitated because of their larger effects and that, on a large scale, this study design is more sensitive for detection of cis than trans effects. We showed that when \( h^2_{\text{trait}} \) is greater than 10%, the expected power to detect major cis-eQTLs is greater than 70% in all tissues. In contrast, detection of trans-eQTLs is limited by low statistical power (12%–60%), as a result of their generally smaller genetic effects. For the eQTLs detected at \( p = 0.05 \), we observed tissue-specific differences in the allelic effects of both cis- and trans-eQTLs. In particular, kidney and adrenal tissues showed enrichment of trans-eQTLs with smaller effects (Figure S6). Although these differences are negligible for detection of cis-eQTLs, they appear to be significant for detection of trans-eQTLs.

We found that, when the major eQTL is trans-acting, minor differences in allelic effects are expected to have a significant influence on the statistical power to detect trans-eQTLs (Figure 3B). As a result, at genome-wide significance, where the majority of eQTLs are regulated in trans, kidney and adrenal showed a higher proportion of false positives among the detected eQTLs (FDR 34%–36%) than that observed in LV and fat tissues (FDR ~26%). Our data indicate that trans-eQTLs account for most of the high FDR expected at \( p = 0.05 \) (Figure 4B), and this goes together with the difference in the statistical power for cis- or trans-eQTLs across tissues. This observation has a number of possible explanations. First, technical variability in microarray experiments (e.g., due to quality of tissue samples or inherent homogeneity of hybridization [31]) may account for the different sensitivity to trans-eQTLs across experiments. However, if this is the case, we should not observe a similar extent of variability and \( h^2_{\text{trait}} \) consistent in all tissues. Therefore, we can exclude technical variation as the primary or sole cause of different sensitivity to trans-eQTL effects. Second, actual differences in tissue composition may influence the ability and power to detect trans-eQTLs by linkage and the FDR in this study design. It is plausible that
the increase of biological variability in more complex and heterogeneous tissues—such as adrenal and kidney—or susceptibility to transcription variation in response to environmental stimuli could explain the observed distribution of trans-eQTL effects.

Although the patterns of heritable gene expression were similar across tissues, the fat eQTL dataset showed a set of 20 trans-regulated genes with high allelic effect (>0.30) but low heritability of the eQTL ($h^2_{QTL}$; 0.2; Figure 1). Inspection of the biological function of these genes revealed that they are all highly expressed in skeletal muscle belonging to the sarcomeric skeletal muscle contractile apparatus. Because the peritoneal fat pad sits in close proximity to the retroperitoneal muscle, and there was marked, but highly correlated, intra-strain variability in raw expression values of this set of genes (i.e., low $h^2_{QTL}$), we consider it likely that this cluster of apparent trans-eQTL effects is due to contamination of the fat pad samples with small quantities of skeletal muscle. Apart from this cluster of skeletal muscle genes, there are very few if any trans-eQTL genes from fat or other tissues showing a similar pattern of low $h^2_{QTL}$ and high eQTL allelic effect. This finding suggests that this possibly confounding effect is limited to this cluster of genes and does not have a significant bearing on the other trans linkages identified in our studies. However, it shows the sensitivity of this experimental design to tissue sampling variability, and the value of this type of analysis for demonstrating the integrity of eQTL datasets.

In summary, this is to our knowledge the first report to address the relative sensitivity to detect cis- and trans-eQTL effects based on the observed patterns of heritable gene expression across multiple tissues and that takes into account their thresholds of detection and FDR. Trans-regulated genes showed small genetic effects that affect the power to detect these eQTLs in different tissues. Although variability in gene expression across tissues is well known [31], we believe that the higher FDR levels for trans-eQTL genes in adrenal and kidney may be correlated to high cellular heterogeneity of those tissues, compared to LV or fat. Importantly, tissuespecific features of the trans-regulated genes are not explained by either observed patterns of heritable gene expression or variability of expression levels. When major-effect eQTLs are mapped in trans, their detection may be precluded by overly stringent statistical cut-offs of significance or by adopting conservative filtering approaches based on trait heritability. This is likely to occur to different extents in different tissues, and we showed that it is a function of the allelic effects of trans-eQTLs. Thus, integration of data from multiple sources requires careful consideration and may, in some cases, not be appropriate. For cis-eQTLs, detection is enhanced by their bigger effects, and the identification of shared cis-acting regulators of gene expression between tissues is facilitated. Because of their more complex and polygenic nature, the profiles of trans-acting eQTLs are likely due to tissue-specific regulation, thus limiting the detectable shared trans-eQTLs across experiments [3,11,16]. By examining genome-wide expression and linkage data from four tissues, we provide here a detailed picture of cis- and trans-eQTL features that may help in understanding the genetic regulation of transcription. This study highlights that large-scale identification of trans-acting loci can be strongly influenced by tissue-specific factors, which are likely to be

![Figure 4. FDR in LV, Fat, Kidney, and Adrenal Tissues](image)
key determinants in the feasibility of discovering regulatory gene networks at the level of the organism.

Materials and Methods

RI strains and tissues. The set of 30 RI strains used in this study was produced by inbreeding between members of the F2 generation resulting from the cross of the two highly inbred strains BN.Lx/Cub and SHR/OLA [18], designated here as BN and SHR. Rats were housed in an air-conditioned animal facility and allowed free access to standard laboratory chow and water. All experiments were performed in agreement with the Animal Protection Law of the Czech Republic (311/1997) and were approved by the Ethics Committee of the Institute of Physiology, Czech Academy of Sciences, Prague, Czech Republic. Animals were killed at 6 wk of age. Tissues from four unfasted males of each RI strain were harvested between 9 and 10 a.m., immediately removed, frozen in liquid nitrogen, and stored at −80°C.

Microarray analysis. The original experimental design is discussed in Hubner et al. [11]. In addition to the fat and kidney data described, we also analysed adrenal and LV data [T. J. Atit, M. Pravenecek, N. Hubner, and S. A. Cook, unpublished data] from the same panel of RI strains and experimental design. Gene expression summary values for Affymetrix GeneChip data (Affymetrix, Santa Clara, California, United States) were computed using the Robust Multichip Average algorithm [32]. In this study, we used a total of 480 microarrays: four animals × 30 RI strains × 4 tissues. Each individual tissue experiment consisted of 120 array datasets, which were processed and normalised separately. For fat, kidney, and adrenal tissues, cRNA was labelled and run on RAE 230A Affymetrix GeneChip arrays (number of transcripts 15,923). For LV tissue, cRNA was labelled and run on RAE 230_2 Affymetrix GeneChip arrays (number of transcripts 31,099).

Expression levels were generated and analysed independently for each tissue. The mean (SE) expression levels measured across 30 RI strains were 3.26 (0.01) for LV, 7.23 (0.02) for fat, 6.55 (0.01) for kidney, and 3.94 (0.01) for adrenal. The experimental variation within each tissue was evaluated through the coefficient of variation of gene expression among all transcripts represented in the microarrays. For all transcripts the average coefficient of variation (SE) was 3% (0.0001) for LV and 2% (0.0001) for fat, kidney, and adrenal tissues. The observed range of coefficients of variation was 0.3%–38% for LV, 0.4%–31% for fat, 0.3%–26% for kidney, and 0.8%–32% for adrenal. These estimates indicate that no significant differences were observed in the intra-specific variability of gene expression levels between tissues.

For this analysis we selected probesets that were mapped to the genome by Ensembl [33]. In order to avoid misclassification of cis- and trans-eQTLs, we also removed probesets that mapped to more than one place in the genome, resulting in 13,669 individual transcripts for fat, kidney, and adrenal and 27,168 individual transcripts for LV.

Heritability of gene expression. The narrow-sense heritability of a trait in a set of RI lines was estimated using the method of Hegmann and Possidente [34]. The heritability was calculated using the simplified formula. Chi-square test for homogeneity and Spearman’s correlations were calculated using SPSS 12.0 (SPSS, Chicago, Illinois, United States).

eQTLs. We carried out genome-wide linkage analysis for each expression trait dataset generated in the RI strains in fat, kidney [11], adrenal, and LV tissues, and 1,011 genetic markers using the QTL Reaper program (http://sourceforge.net/projects/qtlreaper). For each transcript we calculated the likelihood ratio statistics for linkage and its empirical genome-wide significance [35] by permutations as previously described [11]. To account for multiple testing of thousands of expression traits, we calculated the FDR [36] and evaluated the expected number of falsely discovered eQTLs at each level of significance. The FDR of the cis- and trans-acting eQTLs was represented by box-plots [37], using the R package UsingR (http://www.math.csi.cuny.edu/UsingR) [38].

Characterisation of eQTLs. In this study, we empirically defined a cis-acting eQTL as having the peak of linkage within 10 Mb of the physical location of the probe set and a trans-acting eQTL as having the peak of linkage on a different chromosome from where the probe set is located. Adjusted additive effects (i.e., QTL allelic effects) and heritability of the eQTLs (\(h^2\)) were calculated for each cis- and trans-eQTL. eQTL effects are an estimate of the absolute change in the transcript abundance that would be produced by substituting a single allele of one type with that of another type in the population. For each eQTL, its allelic effect is calculated as \(\frac{\text{MEAN}_{\text{SHR}} - \text{MEAN}_{\text{BN}}}{2}\), where \(\text{MEAN}_{\text{SHR}}\) and \(\text{MEAN}_{\text{BN}}\) are the means of gene expression levels of all rats that respectively inherited SHR and BN alleles at the marker peak of linkage. The value \(h^2\) is the proportion of phenotypic variance that is attributable to the eQTL [39], and it is calculated from the squared allelic effect and the error variance segregating in the RI strains [40].

Power calculation. We calculated the expected statistical power to detect a major eQTL effect in the RI strains using the R package Rqtl Design (http://www.biostat.ucsf.edu/software.html) [41]. In order to model various trait heritabilities, we considered different genetic and environmental variances within the experimental range observed in the RI strains across four tissues and used those estimates to calculate the power to detect a given additive effect of the eQTL [40]. The threshold used for detecting an eQTL was log of the odds score = 3.3 (i.e., genome-wide \(p = 0.05\) [42]); 30 RI strains were considered. Similarly, we carried out additional power calculations and modelled the number of RI lines (30–200) and several thresholds of significance corresponding to those reported in this study.

ROC curves. For each transcript, the estimated \(h^2\) was tested as a predictor of the presence of an eQTL (as detected by linkage) via ROC curve. ROC curves are plots of the true positive rate (sensitivity) of a diagnostic test against the test’s false positive rate (1 – specificity) for various decision thresholds (i.e., various \(p\)-value cut-offs). The AUC measures the probability of correct classification, i.e., the ability of the trait heritability to correctly classify those eQTLs that will be (or will not be) observed in the linkage study. For a perfect prediction, the AUC will tend to be close to one and the ROC curve would follow the left-hand border and then the top border of the ROC space [25]. Non-parametric estimates of AUC and its SE were carried out using SSPS 12.0.

Supporting Information

Figure S1. Comparison of Heritability of Gene Expression across Different Tissues

Figure S2. Cumulative Frequency Curves of the Distribution of Transcripts Observed with a Given Heritability of Gene Expression in LV, Fat, Kidney, and Adrenal Tissues

Figure S3. Comparison of ROC Curves for Prediction of Existence of a Detectable eQTL by Heritability of Gene Expression in LV, Fat, Kidney, and Adrenal Tissues

Significantly detected eQTLs were defined at several levels of significance (\(p\)-value cut-offs 0.05, 10\(^{-2}\), 10\(^{-3}\), 10\(^{-4}\), and 10\(^{-5}\)) and the relative ROC curves are compared. The vertical axis shows the fraction of major eQTLs that are correctly predicted (sensitivity) by the \(h^2\) whereas the horizontal axis indicates the fraction of major eQTLs that are incorrectly predicted (1 – specificity) by the estimate of the \(h^2\). The closer the ROC curve is to the upper left-hand corner of the graph, the more accurate is the prediction of the eQTL (i.e., higher true positive rate and lower false positive rate). The areas under the ROC curves are reported in Table S1.

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Found at DOI: 10.1371/journal.pgen.0020172.sg003 (3.5 MB PPT).
Various estimates of heritability of gene expression were modelled, and 30 RI strains were considered (see Materials and Methods). Found at DOI: 10.1371/journal.pgen.0021072.s004 (67 KB PPT).

Figure S5. Statistical Power to Detect an eQTL with Allelic Effect 0.1 (i.e., FC 1.15) as a Function of the Number of RI Strains.
Various estimates of heritability of gene expression were modelled, and four biological replicates were considered. Found at DOI: 10.1371/journal.pgen.0021072.s005 (59 KB PPT).

Figure S6. Cumulative Frequency Curves of the Distribution of Trans-eQTLs Detected with Genome-Wide Significance (p < 0.05) with a Given Allelic Effect in LV, Fat, Kidney, and Adrenal Tissues.
We observe an enrichment of trans-eQTLs with small allelic effects in the range 0–0.8 for kidney and adrenal tissues compared with LV and fat. Insert: cumulative frequency curves of the distribution of trans-eQTLs for allelic effects in the range 0–0.16. Found at DOI: 10.1371/journal.pgen.0021072.s006 (57 KB PPT).

Table S1. AUCs for Prediction of Existence of a Detectable eQTL by Heritability of Gene Expression for LV, Fat, Kidney, and Adrenal Tissues.
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